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OPIOID ABUSE PREVENTION AND TREATMENT AMENDMENTS

2018 GENERAL SESSION

26-55-109, Utah Code Annotated 1953

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27	Be it enacted by the Legislature of the state of Utah:
28	Section 1. Section 26-55-109 is enacted to read:
29	26-55-109. Opiate abuse prevention pamphlet.
30	(1) The department shall produce and distribute, in conjunction with the Division of
31	Substance Abuse and Mental Health, a pamphlet about opiates that includes information
32	regarding:
33	(a) the risk of dependency and addiction;
34	(b) methods for proper storage and disposal;
35	(c) alternative options for pain management;
36	(d) the benefits of and ways to obtain naloxone; and
37	(e) resources if the patient believes that the patient has a substance abuse disorder.
38	(2) The pamphlet described in Subsection (1) shall be:
39	(a) evaluated periodically for effectiveness at conveying necessary information and
40	revised accordingly;
41	(b) written in simple and understandable language; and
42	(c) available in English and other languages that the department determines to be
43	appropriate and necessary.
44	Section 2. Section 58-37-7 is amended to read:
45	58-37-7. Labeling and packaging controlled substance.
46	(1) A person licensed pursuant to this act may not distribute a controlled substance
47	unless it is packaged and labeled in compliance with the requirements of Section 305 of the
48	Federal Comprehensive Drug Abuse Prevention and Control Act of 1970.
49	(2) No person except a pharmacist for the purpose of filling a prescription shall alter,
50	deface, or remove any label affixed by the manufacturer.
51	(3) Whenever a pharmacist sells or dispenses any controlled substance on a
52	prescription issued by a practitioner, [he] the pharmacist shall affix to the container in which
53	the substance is sold or dispensed:
54	(a) a label showing the:
55	(i) pharmacy name and address;
56	(ii) serial number; and

02-26-18 3:24 PM

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container with the label attached.

57	(iii) date of initial filling;
58	(b) the prescription number, the name of the patient, or if the patient is an animal, the
59	name of the owner of the animal and the species of the animal;
60	(c) the name of the practitioner by whom the prescription was written;
61	(d) any directions stated on the prescription; and
62	(e) any directions required by rules and regulations promulgated by the department.
63	(4) Whenever a pharmacist sells or dispenses a Schedule II or Schedule III controlled
64	substance that is an opiate, a pharmacist shall affix a warning to the container or the lid for the
65	container in which the substance is sold or dispensed that contains the following text in not less
66	than 14-point font:
67	(a) "Caution: Opioid. Risk of overdose and addiction"; or
68	(b) any other language that is approved by the Department of Health.
69	(5) (a) A pharmacist who sells or dispenses a Schedule II or Schedule III controlled
70	substance that is an opiate shall prominently display at the point of sale the informational
71	pamphlet developed by the Department of Health under Section 26-55-109.
72	(b) The board and the Department of Health shall encourage pharmacists to use the
73	informational pamphlet to engage in patient counseling regarding the risks associated with
74	taking opiates.
75	[(4)] (6) A person may not alter the face or remove any label so long as any of the
76	original contents remain.
77	[(5)] (7) (a) An individual to whom or for whose use any controlled substance has been
78	prescribed, sold, or dispensed by a practitioner and the owner of any animal for which any
79	controlled substance has been prescribed, sold, or dispensed by a veterinarian may lawfully
80	possess it only in the container in which it was delivered to [him] the individual by the person
81	selling or dispensing it.
82	(b) It is a defense to a prosecution under this subsection that the person being

prosecuted produces in court a valid prescription for the controlled substance or the original